

**Amendments to the Claims**

1. (Original) A ceramic molded body, more specifically a roof tile, tile, clinker brick, or a facade wall of oxide-ceramic base material with a capillary structure and with a surface which is self-cleaning upon spraying or sprinkling with water, characterized in that the molded body has a porous oxide-ceramic coating, wherein the coating is photocatalytically active and the photocatalytically active, oxide-ceramic materials include  $\text{TiO}_2$ ,  $\text{Al}_2\text{O}_3$  and  $\text{SiO}_2$ , wherein the  $\text{Al}_2\text{O}_3$  is aluminum oxide C, and has a specific surface area in a range of between  $25 \text{ m}^2/\text{g}$  and  $200 \text{ m}^2/\text{g}$ , preferably between  $40 \text{ m}^2/\text{g}$  and  $150 \text{ m}^2/\text{g}$ , wherein the mean diameter of the pores or the capillaries is in a range of between  $0.1 \text{ }\mu\text{m}$  and  $5 \text{ }\mu\text{m}$ .

2. (Original) A ceramic molded body as set forth in claim 1 characterized in that the pore openings are of an intergranular and/or intragranular nature.

3. (Original) A ceramic molded body as set forth in one of claims 1 and 2 characterized in that the free breathing cross-section of the ceramic molded body is reduced by the applied porous, oxide-ceramic coating by less than 10%, preferably less than 5%, with respect to the free breathing cross-section of an uncoated ceramic molded body.

4. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of claims 1 through 3~~ characterized in that the porous, oxide-ceramic coating is

applied to a depth of 1 mm, preferably to a depth of 2 mm, measured in a vertical direction from the surface of the ceramic molded body, in the ceramic molded body.

5. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the coating has a specific surface area in a range of between 40 m<sup>2</sup>/g and 100 m<sup>2</sup>/g.

6. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the mean layer thickness of the coating is in a range of between 50 nm and 50 µm, preferably 100 nm and 1 µm.

7. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~, characterized in that arranged between the oxide-ceramic base material and the photocatalytically active, porous, oxide-ceramic coating is at least one layer with raised portions, the oxide-ceramic base material has raised portions and/or the photocatalytically active, porous, oxide-ceramic coating is in the form of a layer with raised portions.

8. (Original) A ceramic molded body as set forth in claim 7 characterized in that the raised portions are formed by particulate material fixed to the oxide-ceramic base material.

9. (Original) A ceramic molded body as set forth in claim 8 characterized in that the particulate material is temperature-resistant ground material preferably

selected from the group which consists of ground rock, fire clay, clay, minerals, ceramic powder such as SiC, glass, glass chamotte and mixtures thereof.

10. (Original) A ceramic molded body as set forth in claim 8 or claim 9 characterized in that the size of the particles and/or the raised portions is or are in a range of up to 1500 nm, preferably of between 5 nm and 700 nm, further preferably between 5 nm and 50 nm.

11. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the oxide-ceramic base material includes photocatalytically active, oxide-ceramic materials selected from the group which consists of TiO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub> and mixtures thereof.

12. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the photocatalytically active, oxide-ceramic material has an average particle size in the range of between 5 nm and 100 nm, preferably between 10 nm and 50 nm.

13. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the TiO<sub>2</sub> contained in the photocatalytically active, porous, oxide-ceramic coating and/or in the oxide-ceramic base material is present at least in part and preferably in respect of at least 40% by weight with respect to the total amount of TiO<sub>2</sub>, in the anatase structure.

14. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the  $\text{TiO}_2$  contained in the photocatalytically active, porous, oxide-ceramic coating and/or in the oxide-ceramic base material is present in respect of at least 70% by weight with respect to the total amount of  $\text{TiO}_2$ , in the anatase structure.

15. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the  $\text{TiO}_2$  is present in a mixture comprising between 70 and 100% by weight of anatase and between 30 and 0% by weight of rutile.

16. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the  $\text{TiO}_2$  is present in respect of about 100% by weight in the anatase structure.

17. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of claims 1 through 16~~ characterized in that the contact angle of a 10  $\mu\text{l}$  drop of water on the porous, oxide-ceramic coating without hydrophobic post-coating after 15 hours of irradiation with 1  $\text{mW}/\text{cm}^2$  UV-A black light is less than  $7^\circ$ , preferably less than  $5^\circ$ , further preferably less than  $4^\circ$ .

18. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of claims 1 through 17~~ characterized in that the contact angle of a 10  $\mu\text{l}$  drop of water on the porous, oxide-ceramic coating without hydrophobic post-coating after 15 hours

of irradiation with 1 mW/cm<sup>2</sup> UV-A black light and 30 days in darkness is less than 20°, preferably less than 18°, further preferably less than 14°.

19. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of claims 1 through 18~~ characterized in that the contact angle of a 10 µl drop of water on the porous, oxide-ceramic coating without hydrophobic post-coating after 15 hours of irradiation with 1 mW/cm<sup>2</sup> UV-A black light and 30 days in darkness and renewed irradiation with preferably 1 mW/cm<sup>2</sup> UV-A black light for 3 hours is less than 8°, preferably less than 7°.

20. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of the preceding claims~~ characterized in that the coating has a superhydrophobic surface, wherein the superhydrophobic surface has a contact or edge angle of at least 140° for water.

21. (Original) A ceramic molded body as set forth in claim 20 characterized in that the superhydrophobic surface is provided using one or more compounds having straight-chain or branched-chain aromatic and/or aliphatic hydrocarbon residues with functional groups, wherein the functional groups are selected from amine, thiol, a carboxyl group, alcohol, disulfide, aldehyde, sulfonate, ester, ether or mixtures thereof.

22. (Original) A ceramic molded body as set forth in claim 21 characterized in that the superhydrophobic surface is produced using compounds selected from the group which consists of silicone oil, amine oils, silicone resin, for example

alkylpolysiloxanes, alkoxysiloxanes, alkali metal siliconates, alkaline earth siliconates, silane-siloxane mixtures, amino acids and mixtures thereof.

23. (Original) A ceramic molded body as set forth in claim 20 characterized in that the superhydrophobic surface of the coating is produced using Ormoceres, polysiloxane, alkylsilane and/or fluorosilane, preferably in combination with SiO<sub>2</sub>.

24. (Original) A ceramic molded body as set forth in claim 20 characterized in that the superhydrophobic surface is applied using a solution of alkali metal siliconates in water, wherein alkali metal is selected from the group which consists of lithium, sodium, potassium and mixtures thereof.

25. (Original) A ceramic molded body as set forth in one of claims 20 through 24 characterized in that the superhydrophobic surface has a contact or edge angle of at least 150° for water, preferably at least 160°, still further preferably at least 170°.

26. (Original) A ceramic molded body as set forth in claim 25 characterized in that the solution of alkali metal siliconates in water has a dilution ratio of between 1:100 and 1:600 (by weight/by weight), preferably a dilution ratio of between 1:250 and 1:350 (by weight/by weight).

27. (Original) A ceramic molded body as set forth in one of claims 20 through 26 characterized in that the superhydrophobic surface of the coating has raised portions.

28. (Original) A ceramic molded body as set forth in claim 27 characterized in that the raised portions of the superhydrophobic surface are produced using particulate material.

29. (Original) A ceramic molded body as set forth in claim 20 characterized in that the superhydrophobic surface is applied using a mixture of particles, for example  $\text{SiO}_2$ , and hydrophobising agent, for example fluorosilane.

30. (Currently Amended) A ceramic molded body as set forth in claim 1, ~~one of claims 1 through 29~~ characterized in that the photoefficiency calculated from photocatalytically induced methylene blue breakdown in the case of the porous, oxide-ceramic coating is at least 0.015%, preferably at least 0.02%, further preferably at least 0.03%, still further preferably at least 0.04%.

31. (Original) A process for the production of a coarse-ceramic molded body, more specifically a roof tile, tile, clinker brick or a facade wall of oxide-ceramic base material with a capillary structure and with a surface which is self-cleaning upon spraying or sprinkling with water, wherein the molded body has a photocatalytically active, porous, oxide-ceramic coating which includes the photocatalytically active, oxide-ceramic materials  $\text{TiO}_2$ ,  $\text{Al}_2\text{O}_3$  and  $\text{SiO}_2$ , wherein the  $\text{Al}_2\text{O}_3$  is aluminum oxide C, with a specific surface area in a range of between  $25 \text{ m}^2/\text{g}$  and  $200 \text{ m}^2/\text{g}$ , preferably  $40 \text{ m}^2/\text{g}$  and  $150 \text{ m}^2/\text{g}$ , wherein the mean diameter of the pores or the capillaries is in a range of between  $0.1 \text{ }\mu\text{m}$  and  $5 \text{ }\mu\text{m}$ , and the porous oxide-ceramic coating is disposed on the surface and in the pore openings and the free faces of the capillary structure to

a depth of 2 mm, measured in a vertical direction from the surface of the ceramic molded body, in the coarse-ceramic molded body,

wherein the process includes the following steps:

(a) mixing photocatalytically active, oxide-ceramic powder which includes  $\text{TiO}_2$ ,  $\text{Al}_2\text{O}_3$  and  $\text{SiO}_2$ , wherein the  $\text{Al}_2\text{O}_3$  is aluminum oxide C, and a liquid phase to afford a suspension,

(b) applying the suspension produced in step (a) to the oxide-ceramic base material to produce a layer, and

(c) hardening the layer afforded in step (b) to produce a photocatalytically active, porous, oxide-ceramic coating.

32. (Original) A process as set forth in claim 31 characterized in that at least one layer with raised portions is applied to the oxide-ceramic base material in a preceding step and the suspension produced in step (a) is applied to the oxide-ceramic base material provided with a layer with raised portions and subsequently hardened in step (c).

33. (Original) A process as set forth in one of claims 31 and 32 characterized in that particulate material is additionally added in step (a).

34. (Currently Amended) A process as set forth in claim 33, ~~claim 32 or claim 33~~ characterized in that raised portions are formed by fixing particulate material on the oxide-ceramic base material.



35. (Currently Amended) A process as set forth in claim 33, ~~claim 33 or claim 34~~ characterized in that the particulate material is temperature-resistant ground material preferably selected from the group which consists of ground rock, fire clay, clay, minerals, ceramic powder such as SiC, glass, glass chamotte and mixtures thereof.

36. (Currently Amended) A process as set forth in claim 33, ~~one of claims 33 through 35~~ characterized in that the mean particle size of the particulate material is in a range of up to about 1500 nm, preferably between 5 nm and 700 nm, further preferably between 5 nm and 50 nm.

37. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 36~~ characterized in that polysiloxane is added to the suspension in step (a).

38. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 37~~ characterized in that water or an aqueous or water-bearing medium is used as the liquid phase in step (a).

39. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 38~~ characterized in that the adhesion between the catalytically active coating and the oxide-ceramic base material is improved by a procedure whereby the photocatalytically active, porous, oxide-ceramic coating produced in step (c) is irradiated with laser light or NIR or UV light.

40. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 39~~ characterized in that contained in the oxide-ceramic base material of the molded body are photocatalytically active, oxide-ceramic materials selected from the group which consists of  $\text{TiO}_2$ ,  $\text{Al}_2\text{O}_3$ ,  $\text{SiO}_2$  and mixtures thereof.

41. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 40~~ characterized in that the photocatalytically active, oxide-ceramic powder used in step (a) includes particles in the range of between about 5 nm and about 100 nm, preferably between about 10 nm and about 50 nm.

42. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 41~~ characterized in that the  $\text{TiO}_2$  contained in the photocatalytically active, oxide-ceramic powder and/or in the oxide-ceramic base material is present at least in part and preferably in respect of at least 40% by weight with respect to the total amount of  $\text{TiO}_2$  in the anatase structure.

43. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 42~~ characterized in that the  $\text{TiO}_2$  contained in the photocatalytically active, oxide-ceramic powder and/or in the oxide-ceramic base material is present in respect of between at least 70% and 100% by weight with respect to the total amount of  $\text{TiO}_2$  in the anatase structure.

44. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 43~~ characterized in that the  $\text{TiO}_2$  contained in the photocatalytically active, oxide-ceramic powder and/or in the oxide-ceramic base material is present in a

mixture comprising between 70 and 100% by weight of anatase and between 30 and 0% by weight of rutile.

45. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 44~~ characterized in that the  $\text{TiO}_2$  contained in the photocatalytically active, oxide-ceramic powder and/or in the oxide-ceramic base material is present in respect of about 100% by weight with respect to the total amount of  $\text{TiO}_2$  in the anatase structure.

48. ~~46.~~ (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 45~~ characterized in that the layer produced in step (b) is hardened in step (c) by drying at a temperature of up to  $300^\circ\text{C}$  and/or by calcining at a temperature of more than  $300^\circ\text{C}$  to  $1100^\circ\text{C}$ .

47. (Original) A process as set forth in claim 46 characterized in that the layer produced in step (b) is at least partially pre-dried prior to the calcining operation in step (c) by evaporation of the liquid phase.

48. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 47~~ characterized in that the coating hardened in step (c) is hydrophobised or superhydrophobised to provide a hydrophobic surface.

49. (Currently Amended) A process as set forth in claim 31, ~~one of claims 31 through 48~~ characterized in that a hydrophobising agent is additionally added in step

(a) and the coating produced in step (b) is hardened in step (c) by drying at a temperature of up to 300°C.

50. (Original) A process as set forth in claim 48 characterized in that an inorganic-organic hybrid molecule, preferably a polysiloxane solution or an alkali metal or alkaline earth siliconate solution is used for hydrophobisation.

51. (Original) A process as set forth in claim 48 characterized in that the superhydrophobic surface is provided using one or more compounds having straight-chain or branched-chain aromatic and/or aliphatic hydrocarbon residues with functional groups, wherein the functional groups are selected from amine, thiol, a carboxyl group, alcohol, disulfide, aldehyde, sulfonate, ester, ether or mixtures thereof.

52. (Original) A process as set forth in claim 51 characterized in that the superhydrophobic surface is produced using compounds selected from the group which consists of silicone oil, amine oils, silicone resin, for example alkylpolysiloxanes, alkoxysiloxanes, alkali metal siliconates, alkaline earth siliconate, silane-siloxane mixtures, amino acids and mixtures thereof.

53. (Original) A process as set forth in claim 48 characterized in that the superhydrophobic surface of the coating is produced using Ormoceres, polysiloxane, alkylsilane and/or fluorosilane, preferably in a mixture with SiO<sub>2</sub>.

54. (Original) A process as set forth in one of claims 48 through 53 characterized in that particulate material is added to produce a superhydrophobic surface with raised portions in the hydrophobisation operation.